Attachment #7

The purpose of this study was to assess the acute dermal toxicity of the PMN substance. An acute dermal toxicity study was conducted in accordance with OECD Guideline 402. The PMN substance was applied to the skin of rats(5/sex) for 24 hours under a semi-occlusive dressing. The dose was 2000 mg/kg for all animals.

No animals died during the 14 day study. Body weights, clinical signs and gross pathology were no affected by treatment. The LD50 >2000 mg/kg.